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Advances in the systematics of the spider genus *Troglohyphantes* (Araneae, Linyphiidae)

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1 **Advances in the systematics of the spider genus *Troglohyphantes* (Araneae,**
2 **Linyphiidae)**

3

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13 **Running title:** Advances in *Troglohyphantes* systematics

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22 **ABSTRACT**

23 With 128 described species and 5 subspecies, the spider genus *Troglohyphantes* (Araneae,
24 Linyphiidae) is a remarkable example of species diversification in the subterranean environment. In
25 this paper, we conducted a systematic revision of the *Troglohyphantes* species of the Italian Alps,
26 with a special focus on the *Lucifuga* complex, including the description of two new species (*T.*
27 *lucifer* n. sp. and *T. apenninicus* n. sp). In addition, we provided new diagnostic drawings of the
28 holotype of *T. henroti* (*Henroti* complex) and established three new synonymies within the genus.
29 The molecular analysis of the animal DNA barcode confirms the validity of this method of
30 identification of the Alpine *Troglohyphantes* and provides additional support for the morphology-
31 based species complexes. Finally, we revised the known distribution range of additional
32 *Troglohyphantes* species, as well as other poorly known alpine cave-dwelling spiders.

33

34 **Keywords:** cave-dwelling fauna, endemism, taxonomy, Italian spiders, species complexes, DNA
35 barcoding

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94 INTRODUCTION

95 The linyphiid spider genus *Troglohyphantes* Joseph, 1881 is presently known to include 128 species
96 and 5 subspecies (World Spider Catalog, 2016), which are generally found in a variety of habitats
97 such as caves, mines, soil litter, rocky debris, and other moistly and shaded retreats (Fage, 1919;
98 Deeleman-Reinhold, 1978; Isaia et al., 2011; Isaia, Lana, & Pantini, 2010). The genus is primarily
99 distributed in the European mountain range, from the Cantabric Mountains in the West to the
100 Balkans and Caucasus in the East. Four additional species are found in Northern Africa (Atlas) and
101 two in the Canary Islands (World Spider Catalog, 2016). The systematics and the distribution of the
102 genus has been studied in a variety of works, with major focus on the Balkan peninsula (Deeleman-
103 Rehinold, 1978), the Pyrenees (Fage, 1919; 1931) and the Alps (Deeleman-Rehinold, 1978; Isaia et
104 al., 2011; Isaia & Pantini, 2010; Pesarini, 1988a, 1988b, 1989, 2001). In general, *Troglohyphantes*
105 species are rare and show narrow distributions. In several cases, they have been reported just from a
106 single or few localities.

107 Because of the high speciosity of this genus, several authors have proposed to assemble the
108 different species in groups based on either phenetic grounds—overall similarity—or authoritative,
109 non-quantitative, phylogenetic hypotheses. The first classification was proposed by Fage (1919)
110 who sorted 21 species—mainly pyrenaic and alpine—into 5 “*Groupes*” (I–V). Diagnoses were
111 based on morphological characters, mostly pertaining to the morphology of male palps and
112 epygines. In 1978 Deeleman-Reinhold revisited Fage’s classification and included 101 species—96
113 of which are presently valid species—which were classified in three series (A, B and C) according
114 to the epyginal morphology. Each series was further subdivided into 12 groups named after the
115 more representative species and based on male palp morphology. However, especially in series B,
116 boundaries between groups remained vague, sometimes including species of difficult placement. In
117 spite of that, most of the authors describing new *Troglohyphantes* species after 1978, classified
118 them following Deeleman-Reinholds’s criteria.

119 A preliminary attempt to classify Italian species was proposed by Thaler (1967) and Brignoli

(1971). Stemming from these early works, Pesarini (2001) retrieved Deeleman-Reinhold's classification and used it as a baseline to sort the Italian species into 11 "*Complexes*", partly overlapping with the extant classifications. Because of the high diversity of the genus in Italy, Pesarini (2001) further created some specific complexes for the Italian fauna, which included Italian species only.

When considering these three classifications and the work of further authors, more than 80% (109 out of 132) of the genus diversity is indeed classified within at least one of the available diagnostic criteria. The highest number of species is classified according to Deeleman-Reinhold's criteria (99 species, 75%) and covers mostly Alpine and Dinaric species. Fage's classification follows, covering a similar geographic range (88 species, 66%), while Pesarini's mostly focuses on Alpine species (38 species, 28%). Overlaps and geographic coverages of the three classifications are illustrated in Fig. 1 and detailed in Supplementary material Table S1.

However, in the absence of a proper morphological or molecular phylogenetic quantitative evaluation, the delimitations of the species groups remains speculative.

Knowledge of the genus in Italy has grown considerably in the last decades, mainly due to the contributions of Pesarini (1988a, 1988b, 1989, 2001) and studies conducted by our research team (Isaia & Pantini, 2008, 2010; Isaia et al., 2010, 2011; Mammola, Isaia, & Arnedo, 2015; Mammola & Isaia, 2016). However, even from a merely taxonomic standpoint, knowledge of *Troglohyphantes* spiders is far from being exhaustive and it is probable that more species have yet to be described.

One of the most speciose group in Italy is Deeleman-Reinhold's *Orpheus* group, which includes nine species from the Alps, one from the Massif Central and two from the Pyrenees. Pesarini (2001) splitted the *Orpheus* group in two complexes—*Lucifuga* and *Orpheus*. The two complexes differ mainly in the general shape of the *lamella characteristica* and have subtle differences in the epyginal structure. The *Orpheus* complex includes four species that exhibit troglomorphic adaptations, such as depigmentation and eye reduction.

145 In the present study, we focused on the *Lucifuga* complex. We described two new species—*T.*
146 *lucifer* n. sp. and *T. apenninicus* n. sp.—, provided a detailed overview of the remaining species
147 included in the group to facilitate their identification and proposed several nomenclatural changes.
148 Furthermore we provided new data on species belonging to other complexes (*Caporiaccoi*,
149 *Diurnus*, *Henroti*, *Microcymbium*, *Orpheus*, *Polyophtalmus*, *Ruffoi* and *Sordelli*) and refined their
150 distribution ranges, illustrated the holotype of *T. henroti* and provided new faunistic data on rare,
151 stenoendemic cave-dwelling spiders collected during our recent surveys. Finally, we used a DNA
152 barcoding approach to help in species delimitation and facilitate identification of Italian species.

153

154 MATERIAL AND METHODS

155 Molecular methods

156 Sequences for the mitochondrial cytochrome *c* oxidase subunit I gene—hereinafter *cox1*, the
157 Animal DNA barcode—were obtained following the protocols described in Mammola et al. (2015).
158 We were able to sample half of the known diversity of *Troglohyphantes* in Italy (17 out of 37
159 species). For each species considered in the molecular analysis, we have reported the relative DNA
160 code in Supplementary Material Table S2.

161 Sequences were edited and managed using Geneious R9 (Kearse et al., 2012). The alignment of the
162 sequences was trivial, as they showed no evidence of indel mutations.

163 Parsimony analysis of the *cox1* matrix was conducted with TNT v.1.1 (Goloboff, Farris, & Nixon,
164 2008) using 1,000 iterations of Wagner trees, followed by TBR branch swapping, and clade support
165 assessed with 1,000 Jackknife resampling replicates—removal probability 36%. The best
166 partitioning schemes and substitution models were assessed simultaneously with PartitionFinder
167 v.1.0.1 (Lanfear, Calcott, Ho, & Guindon, 2012) under a Bayesian information criterion (BIC).

168 Maximum Likelihood (ML) analysis was conducted in RAxML v.7.4.2 (Stamatakis, 2006). We
169 inferred the best ML tree and bootstrap support, automatically determining a sufficient number of

bootstrap replicates, using the MRE convergence criteria. Bayesian (BI) analysis was conducted in MrBayes v.3.2 (Ronquist et al., 2012) with two independent runs of 2 million generations with four Markov chains (one cold, three heated), sampling every 1,000 generations. The chain convergence (ASDSF), the correct mixing (EES) and the number of generation to discard as burn-in were monitored with Tracer v.1.6 (Rambaut, Suchard, Xie, & Drummond, 2014). The first 25% of trees in each run were discarded as burn-in. In all analyses the Canarian species *Troglohyphantes oromii* (Ribera & Blasco, 1986) was used as an outgroup to root the trees.

The ability of the animal barcode to identify species in Italian *Troglohyphantes* was investigated by means of distance metrics (Meier, Shiyang, Vaidya, & Ng, 2006), as implemented in the R package SPIDER 1.3 (Brown et al., 2012). Genetic distances were corrected using the Kimura 2 parameters model, as widely applied in Barcoding analyses, and using the R package APE 3.4 (Paradis, Claude, & Strimmer, 2004). First, we used the nearest neighbour criterion to assign a query sequence to the same species as its closest sequences in the reference library. We then inferred the threshold values that minimised the identification error rates—ie. false negatives or false positives—by optimising the SPIDER function *thereshOptm* testing threshold divergence values from 0.1 to 15%. The presence of a barcoding gap was visualized by plotting the maximum intraspecific divergence to the smallest interspecific divergence.

Taxonomy

Except otherwise stated, specimens are stored in 75% ethanol at the Museo Civico di Scienze Naturali “E. Caffi” of Bergamo (Italy). Additional materials are stored in:

- i) Marco Isaia’s collection (CI) at Dipartimento di Scienze della Vita e Biologia dei Sistemi, University of Torino, Italy;
- ii) Fulvio Gasparo's private collection (CG);
- iii) the Natural History Museum of Bern (NHMB);
- iv) the Museo Civico di Storia Naturale di Milano (MCSNM);

v) the Muséum National d'Histoire Naturelle de Paris (MNHN).
All specimens were hand collected, except otherwise stated.
We studied materials using a Leica M80 stereoscopic binocular microscope. Illustrations of pedipalps and epigynes were made by Elena Pelizzoli using a *camera lucida*. Paolo Pantini provided the diagrams of the female internal genitalia, using *camera lucida* on cleared epygines. All measurements are in mm. Anatomical terms follow Deeleman-Reinhold (1978).
The summary bibliography, the relevant literature and the synonyms for each of the species here presented refers to the World Spider Catalog (2016).
The toponomastics and classification of the different sectors and sub-sectors of the Alps follows the standard partition of the alpine chain (SOIUSA; Marazzi, 2005). Whenever appropriate/applicable, the speleological cadastral codes of the caves are given in squared brackets [regional code and number].
The following abbreviations are used in the text: ALE = anterior lateral eyes; AME = anterior median eyes; MSS = *Milieu Souterrain Superficiel* (*sensu* Mammola et al., 2016); PLE = posterior lateral eyes; PME = posterior median eyes; SA = Suprategular apophysis; SSD = Subterranean Sampling Device; SSH = Shallow Subterranean Habitat (*sensu* Culver & Pipan, 2014); TLL = total leg length; TmI = position of trichobothrium on metatarsus of first leg.

213

214 **RESULTS**

215 **Molecular analyses**

The new sequences obtained in the present study are available in GenBank® (KT831559–KT831596; see Supplementary materials Tables S2).
We generated 48 sequences of 676 bp of the *cox1* DNA barcode, corresponding to 43 unique haplotypes. The mean interspecific genetic divergence (K2P) across the Italian *Troglyphantes* individuals was 16.8% (sd=0.023). The comparison of the minimum interspecific divergences, with the maximum intraspecific divergences revealed the existence of a barcoding gap (Supplementary

Material Fig. S1), with an optimum threshold divergence estimated between 7 and 7.6%. The nearest neighbour criterion resulted in a 100% identification success in species represented by more than one sequence.

The parsimony analysis yielded three trees of 1132 steps. Partitionfinder selected the simple codon—i.e. two partitions 1st and 2nd codon partitions together and 3rd partition apart—as the best partition scheme. The preferred evolutionary model for the 1st + 2nd codon partition was HKY+I+G and for the 3rd codon partition the TrN+G—a GTR was implemented in MrBayes instead. We used unlinked GTR+G models for the RAxML analysis. The results of the analyses conducted under the different phylogenetic inference methods are summarized in Fig. 2. All species showed exclusive haplotypes that formed supported clades. As expected by using a single, highly variable gene, deep relationships were poorly supported and some differed across methods. However, all complexes proposed by Pesarini (2001) were recovered and mostly supported as monophyletic, except for the *Microcymbium* complex, which was recovered as paraphyletic with regards to the *Caporiaccoi*, *Sordellii* and *Henroti* complexes.

TAXONOMIC ACCOUNT

Lucifuga complex

The *Lucifuga* complex so far comprises seven species (Pesarini, 2001, Isaia and Pantini, 2010): *Troglohyphantes albopictus* Pesarini, 1989, restricted to Colli Euganei and Colli Berici—formally outside of the alpine chain, a few km south of the Prealps of Veneto; *T. aldae* Pesarini, 2001 only recorded in the type locality—Asiago plateau, Prealps of Veneto; *T. lucifuga* (Simon, 1884)—from Lanzo Valley to Tessin and Wallis; *T. pluto* di Caporiacco, 1938, endemic to the Corsaglia Valley and high Tanaro Valley—Ligurian Alps; *T. sarae* Pesarini, 2011, endemic to Val d'Aosta; *T. sciakyi* Pesarini, 1989, endemic to the Central Lombard Prealps; *T. subalpinus* Thaler, 1967, restricted to Northern Tyrol and Lower Austria (Fig. 3). Together with the species of the *Orpheus* complex, all species included in the *Lucifuga* complex belong to the *Orpheus* group *sensu* Deeleman-Reinhold

(1978). With *T. subalpinus* (Austria) and *T. lucifuga* (Italy and Switzerland) as the only exceptions, all species included in this complex are Italian endemic.

The complex is characterized by the peculiar dorso-flattened *lamella characteristic*, with the two branches (*sensu* Deeleman-Rehino, 1978) fused for most of their length. In this complex, the *lamella characteristic* is not entirely visible from a lateral view and requires detachment—or at least bulb expansion. The scape of the epygine is always pedunculated —“*palette*” *sensu* Fage, 1919. All the species belonging to this group present an abdominal pattern, which is rather reduced in *T. pluto*.

Here we provided comparative plates illustrating the diversity of the species belonging to this complex (Figs. 4, 5, 6, 7, 8, 9). Moreover, we described two new species and proposed two synonymies within the complex. Accordingly, the complex still includes seven species: *T. albopictus*, *T. apenninicus* n. sp., *T. lucifer* n. sp., *T. lucifuga*, *T. pluto*, *T. sciakyi* and *T. subalpinus*.

***Troglohyphantes albopictus* Pesarini, 1989 (= *T. aldae* Pesarini 2001, new synonymy)**

Figs. 4.1, 5.1, 6.1, 6.2, 7.1, 8.1, 9.1

Material examined

Italy, Veneto, Province of Padova: Rovolon, Colli Euganei, 22.IV.1988, leg. Zanon (male holotype of *T. albopictus*; MCSNM). Province of Vicenza: Valle Scalona (Altopiano di Asiago), in military bunkers, 30.V.1989, leg. Comotti e Baldan (male holotype of *T. aldae*; MCSNM); Mt. Grappa, 10.X.1969 Buche leg. 1♀ 2juv; same locality, 28.IX.1977, Buche leg. 2♂♂; Recoaro, Monti Lessini, Fongara, m 850, 28.IV.1984, Thaler leg. 1♂ 1♀ 1juv; Campodalbero, m 1200, 27.IV.1984 Thaler leg. 2♂♂ 3♀♀ 1juv; Nanto, Colli Berici, 25.IV.1985, Thaler leg. 1♂ 1♀; Schio, Sant'Antonio, Pian delle Fugazze, Rifugio Balasso (1000m), 10.X.1969, 1♂. Trentino Alto Adige, Province of Trento: Borgo Valsugana, Val di Sella, m 900-1200, 30.IX.1977, 2♂♂ 3♀♀ 3juv

Notes

The examination of the holotype of *Troglohyphantes albopictus*, the material from other localities and the holotype of *T. aldae*, allowed the comparison of *lamella characteristic* (Figs. 6.1, 6.2), suggesting that *T. aldae* Pesarini 2001 is a junior synonym of *T. albopictus* Pesarini 1989.

276 Accordingly, the range of distribution of *T. albopictus* now comprises Colli Euganei, Colli Berici
277 and the Prealpi Venete—Asiago plateau.

278

279 ***Troglohyphantes apenninicus* Isaia, Mammola & Pantini new species**

280 Figs. 4.2, 5.2, 6.3, 7.2, 8.2, 9.2, 10

281 *Type series*

282 **Holotype.** Italy, Toscana, Province of Pistoia: Abetone, 10.66270 E 44.14500 N, 15.X.1975, Thaler leg. 1♂.

283 **Paratypes.** Italy, Toscana, Province of Massa: Apuane Alps, Mount Sumbra, 10.27900 E 44.07800 N, 04.VI.1988,
284 Thaler leg. 2 juv Province of Pistoia: Abetone, 10.662700 E 44.14500 N, 15.X.1975, Thaler leg. 4♂♂ 6♀♀ 6juv

285

286 *Diagnosis*

287 Males of *Troglohyphantes apenninicus* are primarily distinguished from other species of
288 *Troglohyphantes* by the shape of the *lamella characteristic* —better viewed if extracted (Fig.
289 6.3),—flattened dorso-ventrally, with the external branch ending with a sharp tooth pointed
290 upwards, almost parallel to the longer axis of the external branch. In comparison with the sister
291 species, in *T. albopictus* (Fig. 6.1), *T. lucifer* n.sp. (Fig. 6.4), and *T. lucifuga* (Figs. 6.5, 6.6) the
292 external branch tapers in a subtriangular apex, bent at the top towards the internal branch.
293 Conversely in *T. apenninicus* (Fig. 6.3), *T. sciaky* (Fig. 6.8), *T. pluto* (Fig. 6.7) and *T. subalpinus*
294 (Fig. 6.9) the apex is not bent, being smaller in *T. subalpinus*, pointed and slender in *T. pluto* and
295 enlarged at its base in *T. sciaky*. The shape of the cymbium (Fig. 5) is also diagnostic, ending
296 proximally with three stout apophysis: the internal apophysis is similar to *T. albopictus* (Fig. 5.1)
297 but stouter, the median apophysis is subtriangular—smaller and more pointed than *T. albopictus* and
298 bigger than the other species of the complex,—and the external apophysis is similar to *T.*
299 *albopictus*.

300 Females are best diagnosed by the epigynum viewed ventrally, by combining the shape of the
301 scape—rhomboidal—with the margins of the pedunculated part—curved,—although differences

across species of the *Lucifuga* complex can be very subtle (Fig. 7). In *T. apenninicus* n.sp. the epyginal plate forms a rhomboidal scape, narrowed at its base and enlarged distally (Figs. 7.2, 10.3). In comparison with other species, the scape is almost round in *T. albopictus* (Fig. 7.1) and trapeizoidal—more enlarged at the base—in *T. pluto* (Fig. 7.5) and *T. sciaky* (Fig. 7.6). The margins of the proximal part of the scape—connecting the scape to the upper part of the epygyne—are curved, unlike the parallel margins found in *T. albopictus* (Fig. 7.1), *T. lucifer* n. sp. (Fig. 7.3) and *T. lucifuga* (Fig. 7.4). Viewed laterally, the scape appears similar to *T. sciaky* (Fig. 8.6), more arched than in the other species of the complex—especially *T. albopictus* (Fig. 8.1), *T. lucifer* n. sp. (Fig. 8.3) and *T. lucifuga* (Fig. 8.4). The identification of females in absence of males remains doubtful in most cases.

Description

Male holotype: prosoma 1.37 long, 1.10 wide, yellowish. Thoracic region slightly swollen, yellowish. Cephalic region slightly elevated, interspersed with black bristles between the eyes. Clypeus slightly indented under the eyes, then convex, 0.31 long. Eyes normally developed, with pigment and black margins. AME smallest. PLE as large as PME, ALE slightly larger than PLE. ALE and PLE almost contiguous. PLE –PME distance = 0.03, ALE–AME distance = 0.04, PME–PME distance = 0.06. Eye diameters AME 0.06, PME and PLE 0.07, ALE 0.10. Sternum heart-shaped, brownish with flimsy darkened anterior edges. Chelicerae 0.65 long, brownish, with ca. 30 lateral stridulatory ridges and armed with three teeth on the anterior side of the chelicerae, and five small, aligned teeth on the internal side. Legs brownish, uniform in colour. Leg I: femur 2.88, patella 0.34, tibia 3.13, metatarsus 2.66, tarsus 1.66, TLL 10.66; leg II: femur 2.69, patella 0.31, tibia 2.72, metatarsus 2.50, tarsus 1.44, TLL 9.66; leg III: femur 2.03, patella 0.31, other articles missing; leg IV: femur 2.81, patella 0.31, tibia 2.69, metatarsus 2.50, tarsus 1.38, TLL 9.69. Abdomen 1.90 long, 1.35 wide; greyish with a dark pattern (Fig. 10.2). Palp (Fig. 10.1): femur 1.28, patella 0.38, tibia 0.31, total palpal length 1.97. Cymbium faintly convex, roughly rectangular when

328 seen from above, ending proximally with three stout apophysis, the external and the medial
329 subtriangular, the internal subrectangular (Fig. 5.2). Posterior part of paracymbium subtriangular, the
330 apical part gradually narrowed anteriorly. *Lamella characteristic* flattened dorso-ventrally, better
331 visible after extraction (Fig. 6.3). External and internal branches fused over most of their length. The
332 internal branch is attached to the radix, made up of two lobes, the smaller branch is short and rounded
333 and the longer one is enlarged distally and separated from the external branch by a depression.
334 Fickert's gland absent. External branch of the lamella, ending with a sharp tooth, darkened at its end,
335 pointed upwards, almost parallel to the longer axis of the external branch. Distal suprategular
336 apophysis (median apophysis *sensu* Deelman-Reinhold, 1978) directed upwards, with a sharp end.
337 Tip of the embolus spiculate. Spination (Tibia and Metatarsus III absent): femur I–II with one dorsal
338 spine; femur III–IV with no spine. Patella I–IV with one dorsal spine. Tibia I with two dorsal, two
339 prolateral, two retrolateral and two ventral spines. Tibia II with two dorsal, two retrolateral spines,
340 one ventral and one prolateral; Tibia III absent; tibia IV with two dorsal, one retrolateral and one
341 prolateral spine. Metatarsus I, II, IV with one dorsal spine. Patella of the Palp with one long, curved
342 spine. TmI: 0.21. Trichobothrium on Mt IV absent.

343

344 *Female* (paratype from same locality as holotype): prosoma yellowish, 1.31 long, 1.06 wide. Cephalic
345 region grey-yellowish. Carapace, ocular area, clypeus, and sternum similar to the male in all features.
346 Clypeus 0.25 long, chelicerae 0.50 long. Anterior margin of the chelicerae armed with three teeth on
347 the anterior side of the chelicerae, and five small, aligned teeth on the internal side. PLE–PME
348 distance = 0.03, ALE–AME distance = 0.04, PME–PME distance = 0.06, AME–AME almost
349 contiguous, ALE–PLE contiguous. Eye diameters: AME 0.06, PME 0.07, ALE 0.10, PLE 0.09.
350 Abdomen greyish, 1.53 long, 1.18 wide, with a dark pattern. Leg I: femur 2.69, patella 0.38, tibia
351 2.97, metatarsus 2.22, tarsus 1.41, TLL 9.66; leg II: femur 2.47, patella 0.34, tibia 2.25, metatarsus
352 2.06, tarsus 0.91, TLL 8.03; leg III: femur 2.06, patella 0.34, tibia 1.56, metatarsus 1.63, tarsus 0.81,
353 TLL 6.50; leg IV: femur 2.59, patella 0.34, tibia 2.50, metatarsus 2.00, tarsus 1.22, TLL 8.66. Female

354 palp: femur 1.13, patella 0.25, tibia 0.75, tarsus 0.44, total palp length 3.50. Spination: Patella of the
355 palp with one dorsal spine, pedipalpal claw present; Tibia of the palp with one dorsal, two prolateral
356 and one retrolateral spine; Tarsus of the palp with three ventral and two prolateral spines, and one
357 retrolateral spine. Femur I–II with one dorsal spine; femur III–IV with no spine. Patella I–IV with
358 one dorsal spine. Tibia I with two dorsal, two prolateral, two retrolateral and two ventral spines. Tibia
359 II with two dorsal, one ventral, two retrolateral and one prolateral spine; Tibia III with two dorsal,
360 one prolateral and one retrolateral spine; tibia IV with two dorsal, one retrolateral and one prolateral
361 spine. Metatarsus I–IV with one dorsal spine. Position of TmI: 0.21. Trichobothrium on Mt IV absent.
362 Epigynum strongly protruding (Figs. 7, 8). Epyginal plate strongly incised, forming a rhomboidal
363 scape, narrowed at its base, enlarged medially and converging distally. Lateral lobes emerging at the
364 posterior end of the epygine (Figs. 7.2, 10.3). Scape arched from a lateral view (Figs. 8.2, 10.4),
365 covering entirely the inner part of the epygine (“*languette interne*” according to Fage, 1919); stretcher
366 tongue-shaped bent upwards toward the scape, bearing a pitted knob at its end, clearly visible from a
367 ventral point of view. Internal genitalia as in Fig. 9.2.

368

369 *Etymology*

370 The species epithet derives from the Latin *Apenninum*, the Apennine mountain range, in which the
371 type series was collected.

372

373 *Distribution*

374 The species is currently known to occur in two localities of the Tuscan Apennines: Abetone (Pistoiese
375 mountains) and Mount Sumbra (Apuan Alps). The species was collected by Konrad Thaler in epigeal
376 localities. No indications about the habitat were given.

377

378

379 ***Troglohyphantes lucifer* Isaia, Mammola & Pantini new species**

380 Figs. 4.3, 5.3, 6.4, 7.3, 8.3, 9.3, 11, 12

381 *Type series*

382 **Holotype.** Italy, Piemonte, Province of Torino: Roure, Tana del Diavolo [Pi 1591], 7.1220669 E 45.0263401 N,
383 12.IX.2014, Isaia & Mammola leg. 1♂ (CI 2566).

384 **Paratypes.** Italy, Piemonte, Province of Torino: Almese, Viù, Colle del Lys, 7.36146 E 45.17751 N, 9.06.2016, Isaia,
385 Mammola & Palermo leg. 3♂♂ 6♀♀ (CI 2883-2884); Graverè, Balma Fumarella [Pi 1597], 7.034559 E 45.125928 N,
386 13.VI.2014, Mammola & Piano leg. 1♀ (CI 2727).

387

388 *Material examined*

389 Italy, Piemonte, Province of Torino: Almese, Viù, Colle del Lys, 10.X.1972, Thaler leg. 7♂♂ 9♀♀ 6juv; same locality,
390 same data, Thaler leg. 3♂♂ 5♀♀; same locality, same data, Thaler leg. 1♂ 1♀; same locality, same data, Thaler leg. 7♂♂
391 8♀♀ 2juv; same locality, same data, Thaler leg. 4♂ 9♀; same locality (in an abandoned house), 9.VI.2016, Isaia,
392 Mammola & Palermo leg. 1♂ 2♀♀ (CI); same locality (in rocky debris), same data, Isaia, Mammola & Palermo leg. 2♂♂
393 4♀♀ (CI); Giaveno, W Forno, 09.X.1972, Thaler leg. 2♂ 7♀ 4juv; Bruzolo, Seinerà mineshaft, 30.IX.2016, Isaia &
394 Mammola leg. 1♀ (CI); Novalesa, Grotta del Ghiaccio di Bosconero [Pi 1580], 15.VII.2006, Lana E. leg. 1♂; same
395 locality, 18.II.2016, Isaia & Mammola leg. 2♂♂ 6♀♀ (CI); Mezenile, Borna Maggiore del Pugnetto [Pi 1501],
396 17.VI.2006, Isaia leg. 1juv (CI 1033); same locality, 29.I.2010, Isaia leg. 1♀ (CI 1057); Same locality, 17.VII.2015,
397 Mammola & Piano leg. 1♂ 2♀♀ (CI 2725); Mezenile, Tana del Lupo [Pi 1502], 17.XII.14, Isaia & Mammola leg. 2♂♂
398 1♀ (CI 2613); Mezenile, "Cavernetta 5" [Pi], 11.III.2016, Mammola & Isaia leg. 1♀; Mezenile, Pugnetto beech forest,
399 1.VII.2012–1.VII.2013, SSD in MSS 0.60 m deep, Isaia & Piano leg. 4 juv (CI 2381); same locality, same data, SSD in
400 MSS 0.80 m deep, Isaia & Piano leg. 1♀ (CI); Mezenile, Pugnetto beech forest (leaf litter), 12.IX.2013, Isaia leg. 3♀♀
401 (CI 2561).

402

403 *Other material*

404 Italy, Piemonte, Province of Torino: Mezenile, Borna Maggiore del Pugnetto [Pi 1501] (Isaia *et al.*, 2010, 2011 sub *T.*
405 *lucifuga*); Mezenile, Borna Inferiore del Pugnetto [Pi 1502] (Isaia *et al.*, 2010, 2011 sub *T. lucifuga*); Novalesa, Grotta
406 del Ghiaccio di Bosconero [Pi 1580] (Isaia *et al.*, 2010, 2011 sub *T. lucifuga*); Novalesa, Boira dal Farfujet o Balma dei
407 Folletti [Pi 1620] (Arnò & Lana, 2005 sub Linyphiidae indet.; Isaia *et al.*, 2010, 2011 sub *T. lucifuga*).

408

409 *Diagnosis*

410 Males of *Troglohyphantes lucifer* n. sp. are primarily distinguished from other species of
411 *Troglohyphantes* by the shape of the *lamella characteristica*, better viewed if extracted (Fig. 6.4).

412 The new species is close to *T. lucifuga* and other species of the *Lucifuga* complex, from which it is

distinguishable by the presence of three teeth-like apophysis on the *lamella characteristic* (Fig. 6.4), two on the external branch and one on the internal branch. Compared to the other species, the teeth-like apophysis on the external branch are unique to *T. lucifer* n.sp. and are absent in other species of the complex. The shape of the cymbium, ending proximally with three stout apophysis and rounded at the proximal border (Fig. 5.3), is also diagnostic. Compared to other species, the internal apophysis is long and slender, with parallel margins—i.e. almost rectangular if compared to other species within the group,—with a rounded apex. Females are best diagnosed by the epyginum viewed ventrally (Figs. 7.3, 11.3), although differences across species of the *Lucifuga* complex can be very subtle (Figs. 7, 8). Compared to other species, the epyginal plate is strongly incised, forming a trapezoidal scape (Fig. 7.3). The margins of the proximal part of the scape—connecting the scape to the upper part of the epygyne—are parallel, similar to *T. albopictus* (Fig. 7.1), *T. apenninicus* n. sp. (Fig. 7.2) and *T. lucifuga* (Fig. 7.4). Viewed laterally, the scape appears similar to *T. albopictus* (Fig. 8.1) and *T. lucifuga* (Fig. 8.4), and less arched than the other species of the complex. The identification of females in absence of males remains doubtful in most cases—see also diagnosis of *T. apenninicus*.

Description

Male holotype (CI2566): prosoma 1.25 long, 1.12 wide, yellowish. Thoracic region slightly swollen, yellowish with grey shades. Cephalic region elevated, interspersed with black bristles between the eyes, with few small black bristles forming the eye region and continuing backwards in three longitudinal rows converging at the thoracic furrow. One bristle just below AME. Carapace with darker margins. Clypeus slightly indented under the eyes, then convex, 0.59 long. Eyes normally developed, with pigment and black margins. AME smallest. ALE slightly bigger than PME and PLE. ALE and PLE contiguous. PLE–PME distance = 0.04, ALE–AME distance = 0.04, PME–PME distance = 0.04. Eye diameters AME 0.06, PME 0.07, ALE 0.09, PLE 0.07. Sternum heart-shaped, yellowish with flimsy darkened anterior edges. Chelicerae light brownish, 0.62 long, with ca. 30 lateral stridulatory ridges and armed with three anterior teeth. Legs yellowish, uniform in colour. Leg

439 I: femur 3.19, patella 0.72, tibia 3.28, metatarsus 2.91, tarsus 1.72, TLL 11.81; leg II: femur 3.00,
440 patella 0.75, tibia 3.13, metatarsus 2.81, tarsus 1.56, TLL 11.25; leg III: femur 2.19, patella 0.41, tibia
441 1.97, metatarsus 1.72, tarsus 1.09, TLL 7.38; leg IV: femur 2.94, patella 0.66, tibia 2.78, metatarsus
442 2.66, tarsus 1.38, TLL 10.41. Abdomen greyish with faint pattern (Fig. 11.2), 2.03 long, 1.31 wide.
443 Palp (Fig. 10a): femur 0.63, patella 0.19, tibia 0.19. Cymbium faintly convex, roughly rectangular
444 when observed from above, ending proximally with three stout apophysis, rounded at the proximal
445 border (Fig. 5.3).

446 Posterior part of paracymbium subtriangular, apical part gradually narrowed anteriorly (Fig. 11.1).

447 *Lamella characteristic* similar to *T. lucifuga* (Fig. 6.5, see Isaia *et al.*, 2011: p. 132, fig. 2.51a),
448 flattened dorso-ventrally, better visible from a dorsal view after extraction (Fig. 6.4). External and
449 internal branch fused over most of their length. The internal branch attached to the radix, made up of
450 two lobes, the smaller branch short and rounded and the longer one enlarged distally, bearing on the
451 outer margin two characteristic teeth-like apophysis, darkened at their tips—see also the paragraph
452 on the synonymy *T. sarae* = *T. lucifuga*—: one placed medially, sharp and well defined, tapering, and
453 pointing towards the internal branch; the other shorter, smaller and less pointed, placed in the notch
454 between the latter and the distal apex of the external branch of the lamella. Fickert's gland absent.

455 External branch of the lamella, ending with a sharp tooth, darkened at its end, pointed towards the
456 internal branch, nearly perpendicular to the longer axis of the external branch. Suprategular apophysis
457 directed upwards, with a sharp end (Fig. 11.1). Tip of the embolus spiculate. Spination: femur I with
458 two prolateral spines; Femur II–IV with no spine. Patella I–IV with one dorsal spine. Tibia I with one
459 dorsal, two ventral, and one retrolateral spine; Tibia II with one dorsal, one ventral, one retrolateral
460 and one prolateral spine; tibia III with one prolateral, one dorsal and one retrolateral spine; Tibia IV
461 with one dorsal, two prolateral, and three retrolateral spines. Metatarsus I–IV with one dorsal spine.
462 Patella of the palp with one long, curved spine. TmI: 0.2. Trichobothrium on Mt IV absent.

463

464 *Female* (paratype from Pugnetto CI 2727): prosoma 1.23 long, 1.00 wide, slightly darker than male.

465 Cephalic region light-brownish. Carapace, ocular area, clypeus, and sternum are similar in all features
466 to the analogous male body parts. Dark brown sternum with dark margins. Anterior margin of the
467 chelicerae armed with three teeth. Clypeus 0.23 long, chelicerae 0.47 long. PLE–PME distance =
468 0.06, ALE–AME distance = 0.04, PME–PME distance = 0.06, AME–AME distance = 0, ALE–PLE
469 distance = 0. Eye diameters: AME 0.04, PME, AME and PLE = 0.07. Abdomen greyish with black
470 pattern (Fig. 12), 2.5 long, 1.87 wide. Leg I: femur 2.50, patella 0.50, tibia 3.22, metatarsus 2.72,
471 tarsus 1.53, TLL 10.47; leg II: femur 2.66, patella 0.56, tibia 2.81, metatarsus 2.50, tarsus 1.38, TLL
472 9.91; leg III: femur 2.19, patella 0.34, tibia 1.88, metatarsus 1.88, tarsus 0.81, TLL 7.09; leg IV: femur
473 2.72, patella 0.38, tibia 2.50, metatarsus 2.41, tarsus 1.25, TLL 9.25. Female palp: femur 0.66, patella
474 0.13, tibia 0.38, tarsus 0.78, total palp length 1.94. Spination (CI 2381): Femur I with one prolateral
475 spine; Femur II–IV with no spine. Patella I–IV with one dorsal spine. Tibia I–II with two dorsal, one
476 prolateral, two ventral and three retrolateral spines; Tibia III with two dorsal, one prolateral and one
477 retrolateral spine. Tibia IV with two dorsal, two ventral and one retrolateral spine. Metatarsus I–IV
478 with one dorsal spine. Patella of the palp with one dorsal spine, pedipalpal claw present; Tarsus of
479 the palp with four dorsal, three retrolateral and three prolateral spines. TmI: 0.2. Trichobothrium on
480 Mt IV absent.

481 Epigynum strongly protruding, with trapezoidal scape (Fig. 11.3), arched from a lateral view (Fig.
482 11.4). Scape diverging distally, with a short stem. Tips of the lateral lobes visible in normal position
483 (Fig. 11.3). Epyginal plate incised, forming a trapezoidal scape, enlarged medially and converging
484 distally. Lateral lobes emerging at the posterior end of the epygine (Fig. 11.3). Scape arched from a
485 lateral view (Fig. 11.4), covering entirely the inner part of the epygine (“*languette interne*” according
486 to Fage, 1919); stretcher tongue-shaped almost straight, abruptly bent upwards toward the scape,
487 bearing a pitted knob at its end clearly visible from a ventral point of view. Internal genitalia as in
488 Fig. 9.3.

489

490 *Etymology*

491 The species epithet derives from the name of the type locality *Tana del Diavolo*—Devil’s lair. Lucifer
492 is the classical traditional Jewish-Christian name assigned to the Devil after the interpretation of a
493 Bible verse from Isaiah. More precisely, Lucifer is the name of the Devil before the Fall from heaven
494 “*to the depth of the pit*” (Isaiah 14: 15). The epithet also recalls the previous misidentifications of
495 *Troglohyphantes lucifer* with *T. lucifuga*.

496

497 *Distribution*

498 The species is known to be found in several localities of a small sector of the Northern Cottian Alps
499 —Viù, Susa and Chisone Valley.

500

501 *Notes*

502 This species was firstly collected in an unspecified epigean habitat by Konrad Thaler in 1972, in the
503 nearby of Colle del Lys and Giaveno—Cottian Alps. The majority of the recent material listed in this
504 contribution was collected on floors and walls of the twilight zone of natural caves as well as in block
505 fields in beech forests. Further specimens were collected in MSS during recent biospeleological
506 investigations at the hypogean complex of Pugnetto—Mezzenile, Lanzo valley, Graian Alps, North-
507 western Alps. MSS was sampled using Subterranean Sampling Devices (SSD; Domingo-Quero &
508 Alonso-Zarazaga, 2010; López & Oromí, 2010) installed at a depth of 0.40–0.80 m.

509

510 ***Troglohyphantes lucifuga* (Simon, 1884) (= *T. sarae* Pesarini, 2011 new synonymy)**

511 Figs. 4.4, 5.4, 6.5, 6.6, 7.4, 8.4, 9.4

512 *Material examined*

513 Italy, Valle d'Aosta, Province of Aosta: Val di Rhêmes, Vandalettaz, 15.X.2006, Fantoni & De Angelis leg. 1♂ (Paratype
514 of *T. sarae*; MCSNM); Brusson, 09.V.1995, Lana leg. 1♂ 4♀♀ (CI 1055); La Salle, Borna d'la Glace [Ao 2001],
515 09.IX.1995, Lana leg. 5♂♂, 7♀♀ (CI 1062); same locality, 16.IX.2014; Isaia & Mammola leg 3♂♂ (CI 2567); Petosan,
516 21.X.2009, Lana leg. 2♂♂, 4♀♀ (CI 1063); St. Rhemy, Fortino presso St.Rhemy, 07.IX.2008, Lana leg. 1♂ 1♀ (CI

517 1067); Verrogne, Fessura di Verrogne [Ao 2017], 09.IX.1995, Lana leg. 4♂♂ 4♀♀ 1juv (CI 1069); same locality,
 518 16.IX.2014; Isaia & Mammola leg 1♂ (CI 2568); Sarre, Mezz'Abisso [Ao 2071], 26.X.2008, Lana leg. 1♂ 1♀ (CI 1073);
 519 Sarre, Grotta della Soldanella [Ao 2072], 26.X.2008, Lana leg. 1♂ 3♀♀ (CI 1074). Piemonte, Province of Verbania:
 520 Macugnaga, Moraine of Belvedere glacier (Mount Rosa), 24/09–12.X.2014, pitfall trap, Tampucci leg. 1♂ 3♀♀;
 521 Sambughetto, Caverna delle Streghe di Sambughetto [Pi 2051], 01.X.2013, Isaia & Mammola leg. 2♂♂ (CI 2573).
 522 Province of Vercelli: Borgosesia, Buco della Bondaccia [Pi 2505], 27.I.2008, Isaia leg. 1♂ 2♀♀ (CI 1054); Valduggia,
 523 Bell'Ingresso [Pi 2539], 17.VI.2009, Lana leg. 1♂ (CI 1071); Valduggia, Bocc d'la Mocia [Pi 2541], 17.VI.2009, Lana
 524 leg. 1♂ (CI 1070). Province of Novara: Alagna, Alpi Pile, 03.X.1971, Thaler leg. 2♂♂ 3♀♀ 1juv; same locality, same
 525 data, Thaler leg. 1♀; same locality, same data, Thaler leg. 3♂♂ 10juv; Alagna, Quarone (Roccapietra), 03.X.1971 Thaler
 526 leg. 1♀ 6juv; Province of Biella: Biella, Santuario di Oropa, 11.X.1972, Thaler leg. 5♂♂, 3♀♀. Province of Torino:
 527 Brosso, Buca del Ghiaccio della Cavallaria [Pi 1609], 07.XI.2014, Isaia & Mammola leg. 3♂♂ 2♀♀ 3juv (CI 2607);
 528 Sparone, Grotta la Custreta [Pi 1593], 07.XI.2014, Isaia & Mammola leg. 2♂ 3♀ (CI 2610); Ceres, Borna del Servais B,
 529 24.IX.2014, Mammola & Paschetta leg. 1♂ 1♀ 2juv (CI 2571); same locality, 14.X.2009, Isaia & Paschetta leg. 1♂ (CI
 530 1061); same locality, 29.IX.2002, Lana leg. 1♂ 1♀ (CI 1077); Ingria, Torrente Soana, 10.X.1972, Thaler leg. 3♂♂, 5♀♀
 531 1 juv; same locality, same data, Thaler leg. 2♂♂ 5♀♀ 3 juv

532

533 *Notes*

534 The comparison of specimens of *T. lucifuga* from different localities of Val d'Aosta and Piemonte
 535 with type material of *T. sarae* Pesarini 2011—described on specimens from Val di Rhêmes and
 536 Valsavaranche, Val d'Aosta—suggests that the latter is a junior synonym of *T. lucifuga*.
 537 Specifically, the observation of the detached *lamella characteristic* revealed a clear
 538 correspondence between the two species (Figs. 6.5, 6.6). The diagnosis of *T. sarae* was based on the
 539 presence of two small and squat teeth on the medial part of the outer margins of the internal and the
 540 external branches of the lamella (Pesarini, 2011, p. 65, f. 3). By examining the detached lamella
 541 (Fig. 6.5) of different specimens of *T. lucifuga* and comparing them with the type material of *T.*
 542 *sarae* (Fig. 6.6), two weak longitudinal bumps were observed on the dorsal part of the branches of
 543 the internal and external lamella, which determine two apparent discontinuities in the profiles of the
 544 distal margins of the lamella. Such discontinuities, identical in *T. sarae*, appear like small teeth from
 545 a lateral point of view, which may explain Pesarini's wrong diagnosis. On this base, we propose *T.*
 546 *sarae* Pesarini, 2011 as junior synonym of *T. lucifuga* (Simon, 1884).

547 *Troglohyphantes lucifuga* is generally found in the outer part of caves, but also in epigean habitats,
548 such as moist shaded places, deep leaf litter and other SSHs. The species is widely distributed in the
549 North-Western Italian Alps—Val d’Aosta and Northern Piemonte. The presence of this species in
550 Switzerland is testified by historical records (Dresco, 1959; Lessert, 1910; Schenkel, 1933) in
551 Wallis (Bourg Saint Pierre and Zermatt) and Tessin (Frasco).

552

553 ***Troglohyphantes pluto* di Caporiacco, 1938**

554 Figs. 4.5, 5.5, 6.7, 7.5, 8.5, 9.5

555 *Material examined*

556 Italy, Piemonte, Province of Cuneo: Frabosa Sottana, Balma Ghiacciata del Mondolè [Pi 102], 22.IX.2015, Isaia,
557 Mammola & Ladame leg. 5♀♀ 9 juv (CI 2644); Roccaforte Mondovì, Grotta dei Partigiani della Tura [Pi 286], 11.X.2015 Lana
558 leg. 1♀ (CI); same locality, 22.V.2016, Chesta & Lana leg. 1♂ 1♀; same locality, 9.VIII.2016, Lana leg 1♀ (CI); same locality,
559 22.VIII.2016, Lana leg 1♀ (CI); same locality, 18.VIII.2016, Giachino & Lana leg 1♂ 2juv (CI); Roccaforte Mondovì, Grotticella
560 della Tura [Pi n.c.], 12.VIII.2016, Chesta & Lana leg. 1♀ (CI); same locality, 22.VIII.2016, Chesta & Lana leg. 1♂ (CI); Roccaforte
561 Mondovì, Plutonis Antrum [Pi n.c.], 16.VII.2016, Chesta & Lana leg 2♂♂ 2♀♀ (CI); same locality, 11.VI.2016, Chesta & Lana leg.
562 1♂ (CI).

563

564 *Notes*

565 The species presence was previously recorded in four localities in the Corsaglia valley (Isaia et al.,
566 2011). Remarkably, the population of Balma Ghiacciata del Mondolè [Pi 102] is found in the
567 twilight zone of the cave, opening in mount Mondolé at 2,071 m asl—Artesina, Province of Cuneo,
568 Italy. The cave is characterized by a very cool microclimatic condition, sustaining a perennial
569 snowfield near the entrance.

570

571 ***Troglohyphantes sciakyi* Pesarini, 1989**

572

573 Figs. 4.6, 5.6, 6.8, 7.6, 8.6, 9.6

574 *Material examined*

575 Italy, Lombardia, Province of Bergamo: Roncobello, Pozzo del Castello [Lo 1310], 31.XII.1989, Comotti & Baldan leg.
576 2♂♂ 2♀♀ 2juv; Castione della Presolana, Passo della Presolana, Monte Scanapa (1600m), 29.IX.1971, 4♂♂ 4♀♀ 7juv
577

578 *Notes*

579 New records for this rare species, previously known to be found in eight localities (Isaia & Pantini,
580 2010; Pesarini, 1989).

581

582

583 **Caporiaccoi complex**

584 The *Caporiaccoi* complex comprises seven Italian species, of which distributions are mostly
585 centred in the Central Lombardian Prealps: *T. caligatus* Pesarini, 1989, *T. caporiaccoi* Brignoli,
586 1971, *T. comottii* Pesarini, 1989, *T. dominici* Pesarini, 1988, *T. iulianae* Brignoli, 1971, *T. spatulifer*
587 Pesarini, 2001 and *T. zanoni* Pesarini, 1988. *T. caporiaccoi* is the only species in this complex
588 showing troglobiomorphic features (Isaia & Pantini, 2010).

589 The complex is characterized by small species (total length around 3 mm) bearing a well developed
590 dorsal median process on the cymbium. The epigyne, which is very enlarged at the base, has a
591 subtriangular scape (“*clavus*” *sensu* Brignoli, 1971).

592 Despite its affinity with Deeleman-Reinhold’s *Diurnus* group, the complex does not overlap with
593 any of the previous classifications and the species included herein are all Italian endemic.

594

595 ***Troglohyphantes caligatus* Pesarini, 1989**

596 *Material examined*

597
598 Italy, Lombardia, Province of Como: between Lasnigo and Barni, m 600, 1.X.1971, 2♂♂ 6♀♀ 3juv; Grotta Tacchi [Lo
599 2029], Zelbio, 19.V.1985, Comotti & Baldan leg. 6♀♀; Pian del Tivano, Grotta Tacchi, 7.V.2016, Isaia, Mammola,
600 Barzaghi, Manenti & Santinelli leg. 2♂♂ 3♀♀ (CI).

601

602 *Notes*

603 A rare species for which we provide new records collected in Triangolo Lariano—Province of

604 Como. The species was known to be found in Monte San Primo (Triangolo Lariano, Italy) (Pesarini,
605 1989) and Monte Generoso (Switzerland) (Hänggi, 1990).

606

607

608 ***Troglohyphantes dominici* Pesarini, 1988**

609

610 *Material examined*

611 Italy, Lombardia, Province of Bergamo: Colzate, pendici Monte Alben, sopra Baite del Sedernello, m 1300, 13.VI.1990,
612 Valle leg. 1♂ 1♀; Gazzaniga, Valle Platz, m 850, XI.1984 Comotti & Valle leg. 1♂.

613

614 *Notes*

615 New records of this species, of which distribution is centred on the Alps and Prealps of Bergamo.

616

617 ***Troglohyphantes iulianae* Brignoli, 1971**

618 *Material examined*

619 Italy, Liguria, Province of Savona: between Pontinvrea and Giusvalla, m 500, 1.X.1972, 1♂ 5♀♀ 3juv (NHMB).
620 Toscana, Province of Lucca: Castelnuovo, Torrente Turrise Secca, 10.406700 E, 44.108500 N., 16.X.1975, Thaler leg.
621 (NHMB).

622

623

624 *Notes*

625 New records of the presence of this species in the Tuscanian Apennines. Previously observed in a
626 few localities in the Ligurian Alps (Brignoli, 1971; Gasparo, 2001; Isaia et al., 2011) and in the
627 Apuan Alps (Pesarini, 2001).

628

629 ***Troglohyphantes zanoni* Pesarini, 1988**

630

631 *Material examined*

632 Italia, Lombardia, Province of Bergamo: Costa Imagna, Pozzo delle Pozzette [Lo 1372], 25.III.1984, Comotti leg. 1♂
633 1♀; Rota d'Imagna, Tomba dei Polacchi [Lo 1003], 1.X.1987, Comotti & Valle leg. 1♂; Villa d'Ogna, m 550, wood,
634 VII-VIII.1985, Pisoni & Valle leg. 2♂♂; Val Taleggio, between Taleggio and Sottochiesa, m 700, 30.IX.1971, Buche
635 leg. 2♂♂ 2♀♀; Treviglio, Parco del Roccolo, m 155, wood, 3.X – 6.XI.2011, pitfall trap, Leoni leg. 1♂; same locality,
636 6.XI – 15.XII.2011, pitfall trap, Leoni leg. 1♂; same locality, 15.III – 19.IV.2012, Leoni leg. 1♂; same locality, 26.VII –
637 21.X.2012, Leoni leg. 1♀; Province of Lecco: Rongio, Grotta Ferrera [Lo 1502], Barzaghi, Isaia, Mammola &
638 Santinelli leg. 1♂ 4♀♀ (CI).

639

640 *Notes*

641 This epigean species shows a disjunct distribution in the Prealps of Bergamo-Brescia and Colli
642 Euganei, possibly related to the fragmentation of the original forest of the Po plain (*Quercus-*
643 *Carpinetum*) (Isaia & Pantini, 2010). This hypothesis is supported by the findings of new
644 populations in a small patch of residual forest in the Po Plain near Treviglio—Parco del Roccio.
645 Interestingly, there are records of the species presence in the Tomba dei Polacchi cave [Lo 1003],
646 the type locality of *T. caporiaccoi* (*Caporiaccoi* complex). The coexistence of two species of
647 *Troglohyphantes* seems to occur only in distantly related groups, as hypothesized by Deeleman-
648 Reinhold (1978).

649

650 **Diurnus complex**

651 In Italy, the *Diurnus* complex comprises the troglobiomorphic species *T. sbordonii* Brignoli, 1975
652 which is found in several localities in Giulie Alps, Giulie Prealps and Carnic Prealps and *T. juris*
653 Thaler, 1982, endemic to the Carnic Prealps. The species of this complex share the features of the
654 homonym Deeleman-Reinhold's group (including three Slovenian species, see Supplementary
655 Material, Table S1), namely the presence of a remarkable dorsal medial apophysis and the simple
656 structure of the cymbium, lacking the two basal divergent processes.

657

658 ***Troglohyphantes juris* Thaler, 1982**

659 *Material examined*

660

661 Italy, Friuli Venezia Giulia, Province of Pordenone: Montereale Valcellina, inghiottitoio Val di Pai [Fr 469],
662 20.VIII.1987, Comotti leg. 1♂.

663

664 *Notes*

665 A new record of this rare species, previously found in very few localities (Thaler, 1982; Pesarini,
666 1989).

667

668 **Henroti complex**

669 The *Henroti* complex comprises *Troglohyphantes vignai* Brignoli, 1971 (Cottian and Ligurian
670 Alps), *T. nigreaerosae* Brignoli, 1971 (Graian Alps, from Gran Paradiso massif to the Lanzo
671 valleys) and *T. henroti* Dresco, 1956, a French endemic species from Isère and Drôme. The complex
672 entirely overlaps the homonym Deeleman-Reinhold's group. Species within this complex are
673 characterized by a well-developed, simple structure of the external branch of the *lamella*
674 *characteristica* and by the triangular scape of the epygine, wide and enlarged at the base.

675

676 ***Troglohyphantes henroti* Dresco, 1956**

677 Fig. 13

678 *Material examined*

679 France, Drôme Department: Maison Forestiere, Lente, 24 .IX.1947 Negre & Henrot leg. (male holotype; MNHN);
680 Grotté des Feès, Col de la Machine, 02.07.1950 Henrot leg. (female paratype; MNHN).

681

682 *Notes.*

683 The species can be found in a few localities of the municipalities of Presles (Isère Department) and
684 Bouvante (Drôme Department), both within the Auvergne-Rhône-Alpes region (Dresco, 1956). The
685 species is illustrated partially in Dresco (1956) and Deeleman-Reinhold (1978). Given the lack of
686 complete diagnostic drawings, we provided new illustrations of the male holotype (Fig. 13.1) and
687 the female paratype (Figs. 13.2, 13.3), for future comparative aims.

688

689 ***Troglohyphantes vignai* Brignoli, 1971**

690 *Material examined*

691 Italy, Piemonte, Province of Cuneo: Frabosa Soprana, Grotta Beppe Bessone (=lo Zucco) [Pi 3303], 22.V.2015, Isaia &
692 Mammola leg. 1♂, 1♀, 1juv (CI).

693

694 *Notes*

695 The species can be found in Cottian and Ligurian Alps with a remarkable distribution gap in the

696 Maritime Alps. We here provided a new record of the species presence in the Ligurian Alps—
697 subsection Alpi del Marguareis,—that slightly widens the range of the distribution of this species
698 eastwards. It is worth noticing that the district of Alpi del Marguareis shows the highest diversity of
699 *Troglohyphantes* within the Western Alps—*T. konradi*, *T. vignai*, *T. pluto*, *T. pedemontanus*, *T.*
700 *iulianae* and *T. bolognai*.

702 **Microcymbium complex**

703 The *Microcymbium* complex comprises four isolated species, all characterized by very small
704 distribution ranges: *T. microcymbium* Pesarini, 2001 (two caves in Prealps of Bergamo), *T.*
705 *bornensis* Isaia & Pantini, 2008 (Pugnetto cave complex, Graian Alps), *T. lanai* (Fenera massif,
706 Pennine Alps), and *T. cavadinii* Pesarini, 1989 (two caves in the Prealps of Bergamo).

708 ***Troglohyphantes microcymbium* Pesarini, 2001**

709 *Material examined*

710 Italy, Lombardia, Province of Bergamo: Sant’Omobono, Grotta di Nala di Cà Maquila [Lo 1135], 10.VI.2016,
711 Santinelli, Manenti, Barzaghi leg. 2♀♀, 1 juv (CI); Province of Lecco: Mandello del Lario, Grotta I Ching [Lo 5079],
712 20.VIII.2008, Aimar leg. 1♂.

713

714 *Notes*

715 Previously found exclusively in the type locality—Sant’Omobono (BG), Grotta Nala di Ca’
716 Maquila [Lo 1135].

717

718 **Orpheus complex**

719 The *Orpheus* complex comprises four troglobiomorphic species distributed in SW-Alps:

720 *Troglohyphantes bolognai* Brignoli, 1975 (one cave in Ligurian Alps), *T. bonzanoi* Brignoli, 1979
721 (one cave in Ligurian Alps), *T. konradi* Brignoli, 1975 (seven caves in Maritime Alps) and *T.*
722 *pedemontanus* (Gozo, 1908) (three caves in Ligurian Alps). The whole complex is part of

Deeleman-Reinhold's *Orpheus* group, which includes species from the Alps, the Pyrenees, one species from Massif Central and one from Caucasus. The complex gets its name from *T. orpheus* (Simon, 1884), a French endemic species of the departments of Aude, Ariège and Pyrénées-Orientales (Simon, 1929). The distal portion of the *lamella characteristic* is not—or only partially—flattened dorso-ventrally. According to the nomenclature change here provided, the complex now includes three species: *T. bolognai*, *T. konradi* and *T. pedemontanus*.

***Troglohyphantes bolognai* Brignoli, 1975 (= *T. bonzanoi* Brignoli, 1979 new synonymy)**

Material examined

Italy, Liguria, Province of Imperia: Pieve di Teco, Sgarbu du Ventu [Li 619], (type locality of *T. bonzanoi*), 27.XII.2014, Isaia & Mammola leg. 1♀ (CI 2600); same locality, 09.XII.2015, Isaia & Mammola leg. 4♂♂, 5♀♀, 3 juv (CI 2645); Badalucco, Tana Bertrand [Li 104] (type locality of *T. bolognai*), 04.IV.2014, Isaia & Mammola leg. 2♀♀, 5 juv (CI 2564); same locality, 27.XII.2014, Isaia & Mammola leg. 1♂ 1♀ (CI 2609).

Notes

This troglobiomorphic species description was based on one female, collected in the Tana di Bertrand cave [Li 104] in Badalucco, Province of Imperia, Liguria (holotype stored at Museum of Verona, P.M. Brignoli's collection, not examined here). Our collection of topotypic material—including the so far unknown male—allowed the comparison with topotypic males of *T. bonzanoi*, a troglobiomorphic species from Sgarbu du Ventu cave [Li 619] (holotype stored at Museum of Verona, P.M. Brignoli's collection, not examined here), a few kilometres away from the type locality of *T. bolognai*. The *lamella characteristic* of the two species was found to be identical. On this base we propose the synonymy *Troglohyphantes bonzanoi* Brignoli, 1979 = *T. bolognai* Brignoli, 1975. The synonymy is also supported by molecular base (see Fig. 2).

749 ***Troglohyphantes konradi* Brignoli, 1975**

750 *Material examined*

751 France, Alpes Maritime: Brigue, Blockhouses in the nearby of Balconi di Marta, 12.VIII.2016, Beikes & Isaia leg. 1♀ (CI).

752

753 *Notes*

754 This record attest for the first time the presence of this species in France.

755

756 ***Polyophtalmus* complex**

757 The *Polyophtalmus* complex includes *Troglohyphantes fagei*, which is widely distributed in the
758 Eastern Alps, the troglobiomorphic *T. scientificus* Deeleman-Rheinold, 1978, endemic to the Giulie
759 Prealps and Giulie Alps, and *T. poleneci* Wiehle, 1964, doubtfully recorded by Pesarini (2001) in
760 Monte Matajur (Giulie Prealps). The complex gets its name from Deeleman-Reinhold's homonym
761 group, which mostly includes Balkanic species (see Supplementary materials Table S1). The female
762 is characterised by the shape of the scape, squat and enlarged at the base. The male paracymbium
763 bears a peculiar "pocket" (*sensu* Deeleman-Reinhold, 1978) and the cymbium lacks dorsal
764 processes.

765

766 ***Troglohyphantes fagei* Roewer, 1931**

767 *Material examined*

768 Italy, Veneto, Province of Treviso: Susegana, Bus de le Fade [V 1271], m 215, 28.IV.1990, Gasparo leg. 1♂ 1♀ (CG).

769 Friuli Venezia Giulia, Province of Udine: Torreano, Foran di Landri [Fr 46], m 425, 19.V.1993 Gasparo leg. 1♀

770 (Gasparo 1997: 20, sub *Troglohyphantes* sp.) (CG).

771

772 *Notes*

773 Mostly found in epigeal habitats in the Austrian and Eastern Italian Alps. Specimens found in Bus

774 de le Fade [V 1271] are characterized by pronounced microphthalmmy.

776 **Ruffoi complex**

777 The *Ruffoi* complex comprises *Troglohyphantes ruffoi* di Caporiacco, 1936 (Prealps of Garda Lake,
 778 Prealps of Vicenza and Colli Berici) and *T. fatalis* Pesarini, 1988 (Prealps of Belluno and Colli
 779 Euganei). Deeleman-Reinhold's included *T. ruffoi* in the *Polyophtalmus* group, but according to
 780 Pesarini (2001), *T. ruffoi* and *T. fatalis* have to be considered in a separate complex. Species
 781 belonging to this group are characterized by the narrowed epyginal scape of the female and by the
 782 presence of two robust spines on the pedipalp tibia of the male. The *lamella characteristic* bears
 783 well-developed apical apophysis.

784

785 ***Troglohyphantes ruffoi* di Caporiacco, 1936**786 *Material examined*

787 Italy, Veneto, Province of Vicenza: Arsiero, Riofreddo Valley, 4.VI – 27.VI.2001, pitfall trap, Busato leg. 1♂ 2♀♀ (CI).

788

789 *Notes*

790 New record of this species, found in both epigeal and hypogean localities of Southern Trentino and
 791 Monti Lessini (Pesarini, 2001).

792

793 **Sordellii complex**

794 The *Sordellii* complex was firstly defined by Thaler (1967) and then retrieved by Pesarini (2001). It
 795 includes four species: *Troglohyphantes sordellii* (Pavesi, 1875) and *T. gestroi* Fage, 1933 (both
 796 endemic to the Prealps of Lugano and Lombardian Prealps), *T. lessinensis* (endemic to Lessini
 797 Mountains), and *T. regalini* Pesarini, 1989 (recorded in a few caves in the area of the Sebino
 798 Bergamasco), the latter being the only troglobiomorphic species within the complex. The complex
 799 only include Italian species showing affinities with Deeleman-Reinhold's *Polyophtalmus* group.

800 This complex is characterized by a sub-rectangular epyginal scape and by the peculiar wrench-
801 shaped *lamella characteristica* and the absence of dorsal processes on the cymbium.

802

803 ***Troglohyphantes gestroi* Fage, 1933**

804 *Material examined*

805 Italy, Lombardia, Province of Brescia: Altopiano di Cariadeghe, Bus del Budrio [Lo 71], 10.III.2016, Santinelli leg.
806 1♀.

807

808

809 ***Troglohyphantes lessinensis* di Caporiacco, 1936**

810 *Material examined*

811 Italy, Veneto, Province of Vicenza: Cornedo Vicentino, Cereda, Grotta del Cameron, 5.VII.1987, Comotti & Baldan leg.
812 1♂ 1♀; Monte di Malo, Bus del Soglio [V 172], 5.VII.1987, 2♀♀ 2juv; same locality, 14.VIII.1988, Comotti & Baldan
813 leg. 2♂♂ 1♀ 1juv

814

815 *Notes*

816 Specimens examined are characterized by a pronounced microphthalmmy.

817

818

819 ***Troglohyphantes regalini* Pesarini, 1989**

820 *Material examined*

821 Italy, Lombardia, Province of Brescia: Marcheno, Caregno, Caia dell'Angili [Lo 405], 1.I.1990, Comotti & Baldan leg.
822 1♂ 4♀♀.

823

824 *Notes*

825 New record of this eyeless species, previously found in a few caves in the Sebino Bergamasco.

826

827 **New data on other subterranean spiders**

828 Given the rarity of these species and the general lack of information available in literature, we here
829 provide new data of occurrence of *Turinyphia clairi* (Simon, 1884) (Linyphiidae) and
830 *Typhlonesticus morisii* (Brignoli, 1975) (Nesticidae) gathered during our recent surveys.

831

832 ***Turinyphia clairi* (Simon, 1884) (Araneae, Linyphiidae)**

833 *Material examined*

834 Italy, Piemonte, Province of Cuneo: Becetto, Pertus dei Draï [Pi 1017], 16.VIII.2015, Isaia & Mammola leg. 1♂ (CI).

835 Liguria, Province of Imperia: Coscio di Arroscia, Garbo di Pian Cavallo [Li 851], 09.I.2016, Isaia & Ladame leg. 1♂

836 (CI); Monte Ceriana, Military blockhouse, 12.VIII.2016, Beikes & Isaia leg. 4♀♀ 1♂ (CI).

837

838 *Notes*

839 Rare species, found in caves and other shaded, damp habitats. In the Western Italian Alps, the

840 species was previously found in only six localities (Arnò & Lana, 2005; Isaia et al., 2011).

841 The Palearctic genus of *Turinyphia* includes four species worldwide: *T. clairi* (Southern Europe), *T.*

842 *maderiana* (Schenkel, 1938) from Madeira, *T. cavernicola* Wunderlich 2008 from Azores and *T.*

843 *yunohamensis* (Bösenberg & Strand, 1906) from Japan. *Turinyphia* species are regarded by Borges

844 and Wunderlich (2008) as relict taxa and palaeoendemic, which survived in mountain areas and

845 islands.

846

847 ***Typhlonesticus morisii* (Brignoli, 1975) (Araneae, Nesticidae)**

848 *Material examined*

849 Italy, Piemonte, Province of Cuneo: Chiusa di Pesio, Grotta Superiore delle Camoscere [Pi 250], 15.VII.1987, Comotti

850 & Baldan leg 1♀; Roccaforte Mondovì, Grotta dei Partigiani della Tura [Pi 286], 22.V.2016, Chesta & Lana leg. 1♀ (CI); same

851 locality, 7.VI.2016, Chesta & Lana leg. 2♀♀ 1juv (CI); same locality, 25.VI.2016, Chesta & Lana leg. 1♀ (CI); same locality,

852 18.VIII.2016, Giachino & Lana leg. 4♂♂ 1♂ (CI); same locality, 22.VIII.2016, Lana leg. 1♂ (CI); Roccaforte Mondovì, Grotticella

853 della Tura [Pi n.c.], 25.VI.2016, Chesta & Lana leg. 1♀; same locality, 12.VIII.2016, Chesta & Lana leg. 2♂♂ 1♀ 2juv (CI).

854 Liguria, Province of Imperia: Triora, Prospetto di Miniera del Monte Grai, 12.VIII.2016, Beikes & Isaia leg. 1♀ (CI). France, Alpes

855 Maritime: Brigue, Balconi di Marta, 12.VIII.2016, Beikes & Isaia leg. 2♂♂ 2♀♀ (CI).

856

857 *Notes*

858 Troglobiomorphic species previously found exclusively in the type locality—Sotterranei del forte A
859 di Vernante, Opera 11 Tetto Ruinas, Vernante. *Typhlonesticus morisii* shares sister species in Spain
860 (*T. obcaecatus*), south-eastern Alps (*T. idriacus*), Montenegro (*T. absoloni*), and Turkey (*T.*
861 *gocmeni*) (Nentwig, Blick, Gloor, Hänggi, & Kropf, 2016; Ribera, Elverici, Kunt, & Özkütük,
862 2014). We here provide several new records from different localities in the Ligurian Alps, including
863 the first French record of this species.

864

865 **DISCUSSION**

866 ***DNA barcode analysis***

867 The large diversity of the genus *Troglohyphantes* in Europe prompted the categorization of the
868 species in different phenetic groups, aiming to achieve a better understanding of the distribution
869 patterns and relationships within the genus. Despite being preliminary, our inferred gene tree, based
870 on the Animal DNA barcode, recovers most of the species complexes concerning the Italian species
871 proposed by Pesarini (2001) (Table 1). These results suggest that the morphological characters used
872 to define those complexes are phylogenetically relevant. Moreover, we have shown that the species
873 can be diagnosed based on their genetic barcode, and that DNA barcoding is an effective tool for
874 species identification in this group. The use of DNA barcodes may greatly help to increase our
875 taxonomic information on the group and better delimit the distribution range of the species by
876 assigning juveniles or by identifying the presence of the species in environmental samples
877 (Thomsen & Willerslev, 2015).

878 The gene tree affinities between lineages and the geographic distribution of the species therein,
879 suggest a pattern of multiple cave colonization processes. This hypothesis is supported by the co-
880 occurrence of representatives of different complexes at a sub-regional scale—sometimes also at the
881 same locality. For example, *T. bornensis* (*Microcymbium* complex), *T. lucifer* n. sp. (*Lucifuga*

882 complex) and *T. nigraerosae* (*Henroti* complex) can be found at short distance from each other
883 within the same valley—Lanzo. Similarly, multiple genetically distant species are present in the
884 Prealps of Bergamo (see Isaia & Pantini, 2010) and in the SW-Alps—Ligurian Alps and Maritime
885 Alps. Therefore, we hypothesize that species communities in *Troglohyphantes* are the result of
886 multiple, independent colonizations, rather than a consequence of local diversification of a single
887 common ancestor. The extant lineages could be interpreted as the result of range contractions and
888 isolation during past climatic changes, followed by population expansions of certain species—i.e.
889 the less troglobiomorphic—during interglacials and other favourable periods. A recent
890 phylogeographic study on *T. vignai* (Mammola et al., 2015) revealed similar patterns, and suggested
891 that long-term climatic changes have been the main cause of the diversification of the genus in the
892 Western Alps.

893 Because of the high diversity of this genus, the rarity of the species and the use of a single,
894 maternally inherited marker, our conclusions based on the molecular analysis should be considered
895 tentative. A more thorough sampling of the species representing the diversity of the genus, and the
896 use of multiple, unlinked DNA markers will be required to provide a well-supported hypothesis of
897 the phylogenetic relationships within the genus. In this regards, a collaborative project aiming to
898 provide a phylogenetic structure of European *Troglohyphantes* and to investigate the origins and
899 diversity of this remarkable group, is currently under way.

900

901 ***Ecology and natural history of the Italian species***

902 *Troglohyphantes* spiders are generally associated with dark and humid habitats, such as caves,
903 artificial subterranean habitats and SSHs. Except for the recent study on the ecological niche of *T.*
904 *vignai* by Mammola and Isaia (2016) and the information provided by Deeleman-Rehinoold (1978),
905 little is known about the specific preferences of *Troglohyphantes* spiders.

906 In this study, we attested for the first time in Italy the presence of *Troglohyphantes* in MSS—i.e., *T.*

907 *lucifer* n.sp. This is not surprising, especially considering that troglobiomorphic spiders have been
908 recently documented in a variety of MSS in Europe (e.g., Arnedo, Oromi, Múrria, Macías-
909 Hernández, & Ribera, 2007; Deltchev, Lazarov, Naumova, & Stoev, 2011; Jiménez-Valverde *et al.*,
910 2015; Laška *et al.*, 2011; Mammola *et al.*, 2016; Nae, 2008; Růžicka & Dolanský, 2016; Růžicka &
911 Klimeš, 2005; Růžicka, Šmilauer, & Mlejnek, 2013; Růžicka & Thaler, 2002). Unfortunately, spider
912 research on the MSS in Italy is still in its infancy. It is most likely that extensive investigations will
913 lead to the discovery of additional hidden biodiversity within the genus.

914 The two new species described in this paper, *T. lucifer* n.sp. and *T. apenninicus* n.sp., can be
915 classified as troglophiles (*sensu* Sket, 2008), having fully developed eyes and abdominal pattern
916 (Figs. 8b, 9b). Troglophile species of *Troglohyphantes* are usually cold adapted (Isaia *et al.*, 2010;
917 Novak *et al.*, 2014), and possess other exaptations to the subterranean conditions—e.g. lucifugous
918 or hygrophilic species.

919 *T. lucifer* n.sp. shows a wide ecological plasticity, being able to colonize a variety of habitats,
920 including the twilight zone of caves and various SSHs—leaf litter, deep soil strata, MSS, rocky
921 accumulations. The species is very similar to—and has been frequently confused with—*T. lucifuga*,
922 both regarding morphology and ecological requirements (Isaia *et al.*, 2010, 2011; Isaia & Pantini,
923 2010). It seems likely that *T. lucifer* represents the ecological vicariant of *T. lucifuga* in the Cottian
924 and Southern Graian Alps. According to our records, in certain localities *T. lucifer* n. sp. is able to
925 coexist with other *Troglohyphantes* spiders. In this regard, Deeleman-Rehinoold (1978) suggested
926 that the co-occurrence of more species of *Troglohyphantes* is rare, and may occur exclusively in
927 phylogenetically distant lineages. The coexistence of two unrelated congeneric species in the same
928 cave was already observed in few caves in Slovenia (Deeleman-Rehinoold, 1978) and in Croatia
929 (Martina Pavlek, personal communication, June 18, 2016). In the Western Italian Alps, the co-
930 occurrence of species of the *Lucifuga* complex and others is documented for *T. lucifuga* - *T.*
931 *nigraerosae* (*Henroti* complex), *T. lucifer* n.sp. - *T. vignai* (*Henroti* complex), *T. lucifer* n.sp. - *T.*
932 *bornensis* (*Microcymbium* complex) and *T. lucifuga* - *T. lanai* (*Microcymbium* complex) (Isaia *et al.*,

2010, 2011; Isaia & Pantini, 2010; this study).

Since the only known records of *T. apenninicus* are uniquely represented by the type series—collected in unspecified epigean habitats in 1975 and 1988 by Konrad Thaler,—the ecology of *T. apenninicus* n.sp. is still unknown. Unfortunately, no information about the habitat are provided on the original labels.

Supplementary Materials

Table S1. *Troglohyphantes* species listed in WSC (2016). For each species, we report the current taxonomic status, the placement—if any—in the phenetic classifications according to Fage (1919), Deeleman-Reinhold (1978) and Pesarini (2001).

Table S2. List of specimens sequenced in this study with voucher information, DNA code and GenBank® access code.

Figure S1. Barcoding analysis of the Italian species of *Troglohyphantes*. Plot of the maximum intraspecific K2P distance against the minimum interspecific K2P distance (left). Values above the 1:1 line indicate the presence of a barcode gap. Plot of false positives (in blue) and false negatives (in red) against genetic divergences, the optimal threshold laid between 7 and 7.6% K2P divergence (right).

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967

968 **Disclosure statement**

969 No potential conflict of interest was reported by the authors.

970

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1174 **Tables**

1175

1176 **Table 1.** The Italian species of *Troglohyphantes* and their affinities to Pesarini’s complexes (2001).

Species	Pesarini’s complexes (2001)
<i>Troglohyphantes bolognai</i> , <i>T. konradi</i> , <i>T. pedemontanus</i>	<i>Orpheus</i>
<i>T. albopictus</i> , <i>T. lucifuga</i> , <i>T. pluto</i> , <i>T. sciakyi</i> , <i>T. apenninicus</i> n.sp., <i>T. lucifer</i> n.sp., <i>T. subalpinus</i>	<i>Lucifuga</i>
<i>T. nigraerosae</i> , <i>T. vignai</i>	<i>Henroti</i>
<i>T. juris</i> , <i>T. sbordonii</i>	<i>Diurnus</i>
<i>T. bornensis</i> , <i>T. lanai</i> , <i>T. microcymbium</i> , <i>T. cavadinii</i>	<i>Microcymbium</i>
<i>T. caligatus</i> , <i>T. caporiaccoi</i> , <i>T. comottii</i> , <i>T. dominici</i> , <i>T. iulianae</i> , <i>T. spatulifer</i> , <i>T. zanoni</i>	<i>Caporiaccoi</i>
<i>T. gestroi</i> , <i>T. lessinensis</i> , <i>T. regalini</i> , <i>T. sordellii</i>	<i>Sordellii</i>
<i>T. fatalis</i> , <i>T. ruffoi</i>	<i>Ruffoi</i>
<i>T. fagei</i> , <i>T. poleneci</i> , <i>T. scientificus</i>	<i>Polyophtalmus</i>
<i>T. exul</i> , <i>T. pavesii</i>	<i>Exul</i>
<i>T. excavatus</i>	<i>Croaticus</i>

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1180 **Figure captions**

1181

1182 **Fig. 1.** Venn diagram illustrating the extant classifications of the species of *Troglohyphantes*
1183 according to Fage (1919), Deeleman-Reinhold (1978) and Pesarini (2001). Numbers in sets indicate
1184 the number of species classified according to the different authors and pie charts refer to the
1185 geographic coverages of the species included therein. Data used to generate the figure are reported
1186 in Supplementary Material, Table S1.

1187

1188 **Fig. 2.** Maximum likelihood tree. Circles on internal nodes denote support values as follow: upper
1189 left: Bayesian posterior probabilities (PP); upper right: maximum likelihood bootstraps (BS),
1190 bottom: parsimony jackknifing (PJ). Filled box: PP>95% or BS>75 or PJ>0.75. Grey boxes: clades
1191 recovered with support values below former thresholds. Empty sectors: clades not recovered. The
1192 tree was rooted using *Troglohyphantes oromii* (Ribera & Blasco, 1986). Pesarini's species groups
1193 recovered as monophyletic in dotted boxes.

1194

1195 **Fig. 3.** Distribution map of the *Lucifuga* complex.

1196

1197 **Fig. 4.** Male pedipalp—*Lucifuga* complex (E = Embolus; LC = Lamella caratteristica; PC =
1198 Paracymbium; SA = Suprategular apophysis; Te = Tip of the embolus). (4.1) *Troglohyphantes*
1199 *albopictus*. Male from Arcugnano (VI) (27.3–12.04.2003 Pantini legit); (4.2) *T. apenninicus* n. sp.
1200 Holotype male (15.X.1975 Thaler legit); (4.3) *T. lucifer* n. sp. Male from Col del Lys, Almese (TO)
1201 (10.X.1972 Thaler legit); (4.4) *T. lucifuga*. Male from Ex-miniera di Cudine, Corio (TO) (Isaia et
1202 al., 2011 fig. 2.51A, modified); (4.5) *T. pluto*. Male from [197 Pi/CN] Abisso Artesinera, Frabosa
1203 Sottana (CN) (Isaia et al., 2011 fig. 2.63A, modified); (4.6) *T. sciakyi*. Male from Camerata
1204 Cornello (BG) (27/8–08.X.2010 Massareo & Zucchelli legit); (4.7) *T. subalpinus*. Male from Styria,
1205 Muraztal S Frein (10.X.1973 Buche legit). Scale: 0.5 mm. Illustration by Elena Pelizzoli.

1206

1207 **Fig. 5.** Cymbium of the right palp—*Lucifuga* complex. (5.1) *Troglohyphantes albopictus*. Male
1208 from Altopiano d'Asiago (VI) (05–06.1989 Comotti & Baldan legit); (5.2) *T. apenninicus* n. sp.
1209 Holotype; (5.3) *T. lucifer* n. sp. Male from Col del Lys, Almese (TO) (10.X.1972 Thaler legit); (5.4)
1210 *T. lucifuga*. Male from [2057 Ao/AO] Grotta VM1, Chatillon (15.V.2006 Lana legit); (5.5) *T. pluto*.
1211 Male from [197 Pi/CN] Abisso Artesinera, Frabosa Sottana (28.VI.2008 Isaia legit); (5.6) *T. sciakyi*.
1212 Male from Alpe Arera (BG) (17.07.2002 Fabbri, Lodovici & Valle legit); (5.7) *T. subalpinus*. Male

1213 from Styria, Muraztal S Frein (10.X.1973 Buche *legit*). Scale: 0.2mm. Illustration by Elena
1214 Pelizzoli.

1215

1216 **Fig. 6.** *Lamella characteristic* (extracted)—*Lucifuga* complex (EB = External branch; IB =
1217 Internal branch; Rad= Radix). (6.1) *Troglohyphantes albopictus*. Male from Arcugnago (VI)
1218 (27.03–12.IV.2003 MSNB *legit*); (6.2) *T. aldae* = *T. albopictus*. Holotype; (6.3) *T. apenninicus* n.
1219 sp. Holotype; (6.4) *T. lucifer* n. sp. Male from Col del Lys, Almese (TO) (10.X.1972 Thaler *legit*);
1220 (6.5) *T. lucifuga*. Male from [2057 Ao/AO] Grotta VM1, Chatillon (15.V.2006 Lana *legit*); (6.6) *T.*
1221 *sarae* = *T. lucifuga*. Holotype; (6.7) *T. pluto*. Male from [197 Pi/CN] Abisso Artesinera, Frabosa
1222 Sottana (28.VI.2008 Isaia *legit*); (6.8) *T. sciakyi*. Male from Alpe Arera (BG) (17.07.2002 Fabbri,
1223 Lodovici & Valle *legit*); (6.9) *T. subalpinus*. Male from Styria, Muraztal S Frein (10.X.1973 Buche
1224 *legit*). Scale: 0.25mm. Illustration by Elena Pelizzoli.

1225

1226 **Fig. 7.** Epigyne, ventral view—*Lucifuga* complex. (7.1) *Troglohyphantes albopictus*. Female from
1227 Monti Berici, Nanto (I) (Thaler *legit*); (7.2) *T. apenninicus* n. sp. Female from S. Abetone
1228 (09.1975–10.1975); (7.3) *T. lucifer* n. sp. Female from Col del Lys, Almese (TO) (10.X.1972 Thaler
1229 *legit*); (7.4) *T. lucifuga*. Female from [art. Pi/TO] *ex miniera di Cudine*, Corio (15.V.2006 Lana
1230 *legit*) (Isaia et al., 2011: fig. 2.51B, modified); (7.5) *T. pluto*. Female from [197 Pi/CN] Abisso
1231 Artesinera, Frabosa Sottana (28.VI.2008 Isaia *legit*) (Isaia et al., 2011: fig. 2.63B, modified); (7.6)
1232 *T. sciakyi*. Female from Mare di Burrasca, Colere (BG) (09 .IX.2015 Massaro, Mazzo, Oneto &
1233 Pantini *legit*); (7.7) *T. subalpinus*. Female from Styria, Muraztal S Frein (10.X.1973 Buche *legit*).
1234 Scale: 0.2mm. Illustration by Elena Pelizzoli.

1235

1236 **Fig. 8.** Epigyne, lateral view—*Lucifuga* complex. (8.1) *Troglohyphantes albopictus*. Female from
1237 Monti Berici, Nanto (I) (Thaler *legit*); (8.2) *T. apenninicus* n. sp. Female from S. Abetone
1238 (09.1975–10.1975); (8.3) *T. lucifer* n. sp. Female from Col del Lys, Almese (TO) (10.X.1972 Thaler
1239 *legit*); (8.4) *T. lucifuga*. Female from [art. Pi/TO] *ex miniera di Cudine*, Corio (15.V.2006 Lana
1240 *legit*) (Isaia et al., 2011: fig. 2.51C, modified); (8.5) *T. pluto*. Female from [197 Pi/CN] Abisso
1241 Artesinera, Frabosa Sottana (28.VI.2008 Isaia *legit*) (Isaia et al., 2011: fig. 2.63C, modified); (8.6)
1242 *T. sciakyi*. Female from Mare di Burrasca, Colere (BG) (09.IX.2015 Massaro, Mazzo, Oneto &
1243 Pantini *legit*); (8.7) *T. subalpinus*. Female from Styria, Muraztal S Frein (10.X.1973 Buche *legit*).
1244 Scale: 0.2mm. Illustration by Elena Pelizzoli.

1245

1246 **Fig. 9.** Diagrams of internal female genitalia, aboral view—*lucifuga* group (s = spermathecae; cg =

1247 copulatory groove; fg = fertilization groove). (9.1) *Troglohyphantes albopictus*. Female from
1248 Campodalbero, Vicenza (V) (Thaler legit); (9.2) *T. appenninicus* n. sp. Female from S. Abetone
1249 (09.1975–10.1975); (9.3) *T. lucifer* n. sp. Female from Col del Lys, Almese (TO) (03.X.1972 Thaler
1250 legit); (9.4) *T. lucifuga*. Female from Alpi Pile, Alagna (NO) (03.X.1971 Thaler legit); (9.5) *T.*
1251 *pluto*. Female from [197 Pi/CN] Abisso Artesinera, Frabosa Sottana (28.VI.2008 Isaia legit) (Isaia
1252 et al., 2011: fig. 2.63C, modified); (9.6) *T. sciakyi*. Female from Mare di Burrasca, Colere (BG)
1253 (09.IX.2015 Massaro, Mazzo, Oneto & Pantini legit). Scale: 0.2mm. Illustration by Paolo Pantini
1254
1255

1256 **Fig. 10.** *Troglohyphantes appenninicus* n. sp. Holotype male and paratype female (15.X.1975 Thaler
1257 legit). (10.1) Retrolateral view of male pedipalp (E = Embolus; LC = Lamella caratteristica; PC =
1258 Paracymbium; SA = Suprategular apophysis; Te = Tip of the embolus). (10.2) Abdominal pattern.
1259 (10.3) Epigyne, ventral view. (10.4) Epigyne, lateral view; Scales: a=0.5 mm; b=1 mm; c–d=0.2
1260 mm. Illustration by Elena Pelizzoli.
1261

1262 **Fig. 11.** *Troglohyphantes lucifer* n. sp. Male and female from Col del Lys, Almese (TO) (10.X.1972
1263 Thaler legit). (11.1) Retrolateral view of male pedipalp (E = Embolus; LC = Lamella
1264 caratteristica; PC = Paracymbium; SA = Suprategular apophysis; Te = Tip of the embolus). (11.2)
1265 Abdominal pattern. (11.3) Epigyne, ventral view. (11.4) Epigyne, lateral view. Scales: a=0.5 mm;
1266 b=1 mm; c–d=0.2 mm. Illustration by Elena Pelizzoli.
1267

1268 **Fig. 12.** *Troglohyphantes lucifer* n. sp. Female from [1502 Pi/TO] Borna inf. del Pugno, Pugno
1269 (TO) (11.III.2016 Isaia, Mammola, Piano, Tomasinelli leg.). (12.1) Abdominal pattern, dorsal view
1270 (12.2–12.4). Live specimen. Photo by Francesco Tomasinelli.
1271

1272 **Fig. 13.** *Troglohyphantes henroti* Dresco, 1956. Holotype male and paratype female (02.VII.1950
1273 Henrot legit). (13.1) Retrolateral view of male pedipalp (E = Embolus; LC = Lamella
1274 caratteristica; PC = Paracymbium; SA = Suprategular apophysis; Te = Tip of the embolus); (13.2)
1275 Epigyne, ventral view. (13.3) Epigyne, lateral view. Scales: 2.1 = 0.5 mm; 2.2–2.3 = 0.2 mm.
1276 Illustration by Elena Pelizzoli.